

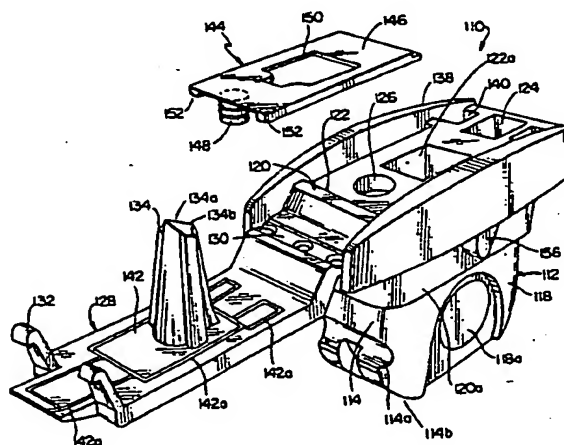
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<p>(21) International Application Number: PCT/US92/04882</p> <p>(22) International Filing Date: 9 June 1992 (09.06.92)</p> <p>(71) Applicant: MODULAR DIAGNOSTIC SYSTEMS, LTD. [US/US]; Route 12, P.O. Box 358, Barneveld, NY 13304 (US).</p> <p>(72) Inventors: MURPHY, Harold, R. ; 209 N. Bellinger Street, Herkimer, NY 13350 (US). DUBOIS, Jeffrey, A. ; R.D. #2, Box 424, Wynn Road, Holland Patent, NY 13350 (US). STRICKLAND, Reid, A. ; R.D. #2, Box 424AA, Wynn Road, Holland Patent, NY 13350 (US). WOOD, Harold, F. ; 4 Deerpath Court, New Hartford, NY 13413 (US).</p>	<p>(74) Agent: HALEY, Barry, L.; Malin, Haley, McHale, DiMaggio & Crosby, One East Broward Boulevard, Suite 1609, Fort Lauderdale, FL 33301 (US).</p> <p>(81) Designated States: AU, BR, CA, JP, KP, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LU, MC, NL, SE).</p> <p>Published <i>With international search report.</i></p>	

(54) Title: CUVETTE FOR AUTOMATED TESTING MACHINE



(57) Abstract

A consumable, non-reuseable cuvette (10) for containing a sample or specimen during an automated test thereof, primarily for medical diagnostic purposes. The cuvette has a single, main reaction chamber that is pre-loaded at the factory with the precise quantity of a particular liquid or dry reagent useful for a specific test. The cover (20) of the cuvette includes an opening (26a) to permit the introduction of a diluent or liquid reagent into the reagent chamber, a manually loaded, recessed sample receiving chamber having a frangible bottom floor (36) and a purge reservoir. A cap (28) is hinged to the cuvette cover and includes a rigid protruding member (34) that pierces the sample or specimen chamber floor when closed by the testing machine, sealing the contents of the cuvette, allowing the sample to be dispensed into the chamber containing the reagent and diluent. The side walls (18) and floor (16) of the reaction chamber include optically transparent windows for radiant energy testing of the reagent before and after the sample is added to the reagent. Because the cap seals the contents of the cuvette, the cuvette is safely disposable after the test is completed.

CUVETTE FOR AUTOMATED TESTING MACHINE

This application is a continuation-in-part application of U.S. Patent Application Serial Number 07/425,346, filed October 13, 1989, which is a continuation application of No. 07/253,383, filed October 3, 1988, a continuation application No. 07/042,795, filed April 27, 1987.

BACKGROUND OF THE INVENTION

1. Field of the Invention

This invention relates to a disposable cuvette for holding a test sample or specimen, reagent, and diluent while performing a test in an automated machine, primarily for medical diagnostic purposes. The disposable cuvette can be prepackaged with a dry reagent to which a diluent is added during a test or with a liquid reagent thereby eliminating the need for the addition of a diluent in an automated test machine.

2. Description of the Prior Art

Cuvettes utilized for manual or automated medical testing are well known. In general, a procedure is established to measure the emergent wavelength of radiant energy absorbed by a sample under analysis. In a manual test, typically a technician loads a cuvette with a sample, reagent and diluent necessary to accomplish the test, each ingredient being precisely measured. The contents are mixed and the emergent radiation is observed optically or ocularly. Because the kind and precise quantity of reagent and diluent employed are critical to a successful test result, technician time and expertise to prepare and use cuvettes are significant factors to be considered with regard to cost, human error, and emergency diagnosis. Preparation time has also become a significant factor with the introduction of automated medical diagnostic machines. Such machines and their corresponding cuvettes have attempted to reduce preparation time by prepackaging various reagents and/or diluents in a container that itself

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2 ultimately houses a test performed therein. U.S. Patent
3 3,504,376 issued March 30, 1970 to Bednar et al. shows such
4 a system.

5 A significant factor in the use of automated test
6 equipment is that the reagent and the diluent must be mixed
7 prior to the addition of the sample so that the emergent
8 radiation from the reagent-diluent mixture can provide a
9 baseline measurement that is compared to the emergent
10 radiation from the mixture that includes the sample. This
11 factor precludes mixing the sample at the same time the
12 diluent is added to the reagent. In the present invention,
13 the sample or specimen is manually loaded in a separate
14 chamber in the cuvette where it remains until after the
15 reagent and diluent have been mixed and the baseline
16 measurement of the reagent-diluent taken. The use of
17 automated diagnostic test equipment still requires that all
18 ingredients necessary for a specific test be precisely
19 measured regardless of whether the cuvette is preloaded at
20 the factory or loaded at the test site.

21 The use of multiple, separated compartments in
22 testing vessels with automated machines is shown in U.S.
23 Patent 3,504,376 issued to Bednar et al on March 31, 1970
24 (cited above); U.S. Patent 4,458,020 issued to Bohn et al.
25 on July 3, 1984; and U.S. Patent 4,473,530 issued to Villa-
26 Real on September 25, 1984. Each vessel shown is a complex
27 in physical structure and requires complex interaction with
28 the test equipment for operation.

29 Disposal of test containers having the residual
30 samples and reagents therein poses a significant
31 environmental waste problem. Washing and reusing a cuvette
32 is not a good practice because the test results could be
33 affected by a poorly washed cuvette. With the present
34 invention, the cuvette remains sealed after the test and is
35 not reusable.

36 As the use of prepackaged cuvettes increases in
37 volume, reducing the cost of the manufacture and factory
38 loading of the cuvette also becomes important.

SUMMARY OF THE INVENTION

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3 A disposable cuvette for performing a
4 predetermined medical diagnostic test of a specimen or
5 sample in an automated machine comprising a hollow body
6 shaped to form a container (having an open top) for use as a
7 single reagent chamber. A cover is sealably attached over
8 the open top of said body, said cover having both an
9 aperture for introducing a diluent into the container body
10 and a specimen receiving chamber with a frangible floor.
11 Attached to the cover is a hinged cap that includes a post,
12 sized and positioned to pierce the floor of the specimen
13 chamber whenever the cap is closed over the cover. A
14 precisely measured quantity of a reagent is loaded into the
15 container at the factory.

16 The specimen receiving chamber is integrally
17 formed within the cover and includes a frangible bottom
18 floor that can be broken open by action of the cap post. The
19 top of the specimen chamber may be covered by a thin hot-
20 stamped film having a central annular opening for receiving
21 the end of a pipette for introduction of the specimen into
22 the specimen chamber.

23 The diluent dispensing aperture in the cover may
24 include depressible flaps closely adjacent to each other but
25 separated by an "X" shaped space. The flaps act as a closure
26 for the body reaction chamber, but are easily opened by a
27 diluent dispensing device.

28 The cover is joined and sonic welded or otherwise
29 bonded about the upper rim of the container body itself.

30 The cuvette is used for testing a sample or
31 specimen as follows. Initially when the cuvette is
32 manufactured and prepared for market, a reagent of
33 predetermined kind and precisely measured quantity for a
34 particular test is placed in the container body. The cover
35 is welded to the container body, but the cap, hinged to the
36 cover, is not closed. A bar coded label identifying the
37 particular diagnostic test to be performed in the cuvette is
38 affixed to the outside wall of the cuvette body. The cuvette

1
2 is then shipped with the cap in the open position to the
3 site where the test is to be performed. A removable
4 adhesive-backed film may be used to cover the top surface of
5 the cover to prevent contamination prior to use.

6 Once at the test site, after the adhesive-backed
7 film is removed from the cover surface, the cuvette is
8 readied for the automated test by the introduction of the
9 specimen into the specimen receiving chamber in the cuvette
10 cover. This loading is done manually by a technician
11 inserting the end tip of a pipette containing the specimen
12 into the annular opening in the top of the specimen
13 receiving chamber in the cuvette cover.

14 The cuvette (now loaded with reagent and sample in
15 a separate chamber) is placed into a circular cuvette holder
16 (carousel) within the automated test machine. The cuvette
17 while in place in the carousel is moved through a series of
18 operations in the machine as follows:

19 1. The carousel moves the cuvette to the bar
20 code reading station. This identifies the test to be
21 performed.

22 2. Diluent addition is performed at the second
23 station. An automated dispensing arm pivots into position
24 into the diluent receiving aperture of the cuvette. As
25 diluent fills the reaction chamber, air is vented through
26 the .005 inch diameter holes until diluent reaches the holes
27 and the resulting surface tensions stops the fluid flow.

28 3. A vibrating action is now started to
29 thoroughly mix the diluent and dry reagent to produce a
30 working unit dose reagent.

31 4. The machine then measures the emergent
32 radiation from the working reagent to obtain a baseline
33 measurement, i.e. reagent absorptivity only.

34 5. The machine then moves the cuvette cap to a
35 closed position over the cuvette cover forcing the post to
36 break through the specimen chamber floor causing the patient
37 sample (urine, serum, plasma) to flow into the chamber
38 containing the reagent and diluent. The cuvette at this

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point is entirely sealed by the cap.

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6. A vibrating action is now started to mix the specimen and reagent.

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7. There is an incubation followed by a short vibration, then the radiant energy absorption test is performed. The difference in absorptivity of the test minus the baseline reading permits calculation of a result based upon a standard value for that test lot which is supplied by the bar code label to the instrument.

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8. The cuvette can now be removed from the machine and safely discarded because the reacted specimen cannot escape from the cuvette once the cap is closed. This feature reduces or limits the release of chemicals and reacted patient samples in the hands of medical office personnel. The cuvette is not reusable.

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In an alternate embodiment the invention may be used wherein a liquid reagent is introduced into the cuvette at the factory. With this embodiment and the replacement of the dry reagent, the addition of a liquid diluent during the machine test is eliminated. In order to prevent leakage of the liquid reagent, an additional gasket/plug is provided on the top cover. This is essentially a planar sheet of a liquid impervious material and includes on one side a plug sized to fit firmly into the aperture in the cover that was previously used for the diluent addition. Also the cover has been modified to eliminate the vents that were required when adding liquid diluent which is used with the dry reagent.

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It is an object of this invention to provide an improved cuvette that reduces the time required to prepare a cuvette for a specimen or sample test in an automated machine.

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It is another object of the invention to provide a cuvette that reduces the potential for human error by including a premeasured reagent and a reagent-diluent chamber of a predetermined volume when filled for a particular diagnostic, medical test that requires a precise reagent/specimen/diluent quantity ratio.

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2 And yet another object of this invention is to
3 provide a disposable, non-reusable cuvette that can be used
4 to safely dispose of the specimen and the reagent after the
5 test is completed.

6 Another object of the invention is to provide a
7 cuvette that is non-complex in manufacture, that is readily
8 pre-loaded with a reagent at the factory, that is easily
9 manipulated by a technician during loading of the diluent
10 and sample, and that is suitable for use in automated
11 diagnostic testing machines.

12 But yet still another object of this invention is
13 to provide a disposable cuvette that can be used with a
14 preloaded liquid reagent at the factory, that does not
15 require the addition of a diluent during the test in an
16 automatic diagnostic testing machine.

17 In accordance with these and other objects which
18 will be apparent hereinafter, the present invention will be
19 described with particular reference to the accompanying
20 drawings.

21 BRIEF DESCRIPTION OF THE DRAWINGS

22 Figure 1 is a perspective view of the present
23 invention with the cap in the open position.

24 Figure 2 is a side elevational view of the
25 invention in cross section, with the cap in the open
26 position.

27 Figure 3 is a side elevational view of the present
28 invention, in cross section, with the cap in the closed
29 position.

30 Figure 4 is a top plan view of the invention with
31 the cap in the open position.

32 Figure 5 is a side elevational view of the
33 invention with the cap in the closed position.

34 Figure 6 is an end elevational view of the
35 invention looking at the end farther from the cap hinge.

36 Figure 7 is an enlarged side elevational view of
37 the invention, in cross section, with the cap in the open
38 position.

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Figure 8 is a top perspective view of an alternate embodiment of the invention that includes (as shown exploded) a gasket plug panel that is affixed to the top of the cover.

Figure 9 shows a bottom perspective view of the alternate embodiment of the invention.

Figure 10 shows yet another embodiment of the invention that is used with a prepackaged dry reagent only.

Figure 11 shows a side elevational cross sectional view of the alternate embodiment construction of the cuvette body and the specimen receiving chamber.

Figure 12 shows a bottom perspective view of the cover used in the alternate embodiment of the invention.

PREFERRED EMBODIMENT OF THE INVENTION

Referring now to the drawings, and in particular Figure 1, the invention is shown generally at 10 comprised of transparent body 12 and a cover 20.

The cuvette body 12 is a vessel or hollow container that includes end walls 14 connected to side walls 18 and bottom wall 16 forming a hollow rectangular box with an open top. The body 12 may be molded as one piece from a clear acrylic material, which is impervious to moisture or atmosphere. The body material is selected to be sufficiently transparent to permit radiant energy absorption testing of a specimen or sample contained within the body which serves as a reaction chamber as will be further discussed. By way of example but not limitation, the dimensions of the cuvette body 12, stated in inches, may be .545 in length, .312 in width, and .545 in height. Of course the dimensions may vary widely without departing from the scope of the invention. It is important that the volume of the reaction chamber be such that the reaction chamber when filled acts to provide the exact volume for diluent added to the cuvette for a predetermined test.

The cuvette 10 also includes a cover 20 that is sonic welded or otherwise bonded to body 12. The cover 20 provides several elements that are important for the

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2 different operational testing stages in the automated
3 machine.

4 As shown in Figures 2, 3, and 4, these elements
5 include the sample or specimen receiving chamber formed by
6 the cover barrier wall 22 having a vertical passage 22a
7 sealed at its lower end by a frangible, thin floor 36 and
8 partially covered at its upper end by a thin film 24 having
9 a circular opening near its center for receiving the end of
10 a pipette. The cover passage 22a in conjunction with the
11 bottom floor 36 form the receiving chamber for serum or
12 sample that is introduced by pipette through the opening in
13 thin film 24.

14 Another important element of cover 20 is a second
15 vertical passage 26a that is partially covered by flaps 26
16 at its upper end. The flaps 26 are flexible and may be
17 opened downwardly by a pipette or other conduit shaped
18 injector to permit the introduction of a diluent into the
19 reaction chamber formed by body 12.

20 The cover 20 also has four .005 inch diameter vent
21 holes 44 that are very important in the operation of the
22 cuvette. The vent hole diameter is sized appropriately so
23 that air is vented to the outside as the diluent flows into
24 the reaction chamber. When the chamber is full, the surface
25 tension created by the .005 inch diameter holes prevents any
26 overflow of diluent through the vent holes 44.

27 The cover 20 includes a cap 28 attached by hinges
28 30. Protruding from one side of cap 28 and substantially
29 perpendicular thereto is a lancet-like post 34 having a
30 "cross" shaped cross section relative to its longitudinal
31 axis. The post 34 is sized in length and positioned relative
32 to hinges 30 such that when cap 28 is closed over cover 20,
33 post 34 will be moved into passage 22a (serum receiving
34 chamber) piercing an opening in floor 36. The floor 36 is
35 connected to the chamber side walls 22a such that it cannot
36 be detached by the action of post 34. (If detached, floor 36
37 might interfere with light passage through the cuvette.)
38 Also attached to cap 28 is a latch 32.

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3 After the serum or sample has been manually added
4 to the receiving chamber 22a, the cuvette is now ready to be
5 loaded into an automated testing machine. Note that the cap
6 28 is in the open position when the cuvette is placed in the
7 test machine. Once in the machine, the cuvette bar code
8 label is read and the test parameters are automatically set.
9 Next the diluent is automatically dispensed into the cuvette
10 and a vibrating mixing action takes place. After the reagent
11 and diluent are fully mixed, a baseline radiant energy
12 absorption reading is taken. After this initial measurement,
13 an interposer with roller bearing mechanism (not shown) in
14 the machine closes the cap 28, and the cuvette, now sealed,
15 is vibrated. As the cap 28 is closed, the post 34 breaks
16 through the sample chamber floor 36, allowing the sample to
17 be mixed with the reagent-diluent mixture, aided by the
18 vibration. With the cap in the closed position, the cuvette
19 is completely sealed such that its contents cannot escape.
20 The cap 28 is fastened by the engagement of latch 32 with a
21 flange on bar 40. The machine can now complete the test on
22 the sample.

23 The cover 20 may be made from high impact styrene
24 and sonic welded to the upper rim of body 12. Hinges 30 and
25 cap 28 may be formed with the cover 20, with hinges 30 being
26 "living" hinges.

27 Figures 5 and 6 show the cuvette 10 after the cap
28 28 is closed. The cuvette, in accordance with the invention
29 described herein, is designed for use in an automated
30 testing machine. The cap 28 includes parallel ridges 28a
31 that project above the upper surface of cap 28 to act as cap
32 stiffeners and a bearing contact surface for mechanically
33 closing the cap.

34 Figure 7 shows the cover 20 and the serum or
35 sample receiving chamber 22a and molded bottom floor 36 with
36 a thinner flash section 36a (formed in an "X" shape) to
37 provide break lines when the post 34 applies pressure as the
38 cap 28 is closed.

The top of the serum chamber 22a is partially

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2 covered by hot stamped film 24. Film 24 has a small opening
3 burned in the center to act as access for the end tip of a
4 pipette containing serum or sample and also to wipe the
5 pipette tip dry as it is removed from the chamber 22a.

6

7 The top surfaces of the diluent access area (flaps
8 26 and adjacent area) are made parallel to the underside
9 surface of the cap 28 that abuts the diluent access
10 perimeter in order to provide a parallel seat for the
11 machine actuated, diluent dispenser tip and achieve a good
12 seal when the cap 28 is closed.

13

14 Figures 1 and 7 show vent holes 44 that aid in the
15 filling of the reagent chamber with diluent by allowing air
16 that would otherwise be trapped in the reaction chamber to
17 escape. Note that the diameter of each vent hole is sized so
18 that diluent will not escape through the vent holes because
19 of the surface tension of the diluent over the vent opening.
20 However, the vent holes are sealed from the ambient air when
21 the cap is closed. After the test has been completed, the
22 cuvette can be safely discarded because the contents are
23 sealed from the ambient surroundings.

24

25 The cuvette and the specific test to be performed
26 therein (dependent upon the kind and quantity of reagent it
27 contains) is identified by labels 46 adhesively attached to
28 each side 38 of the cover. One label on one side would have
29 a machine readable bar code while the other side label would
30 display a letter or number code that is easily identified by
31 the test operator.

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33 Referring now to Figure 8 an alternate embodiment
34 of the invention is shown which is used with a liquid
35 reagent. The liquid reagent is prepackaged in the cuvette at
36 the factory. By using a liquid reagent, it is not necessary
37 to provide the liquid diluent required with a dry reagent in
38 the automated test device when performing a test. Therefore,
in general, the cuvette 110 is modified (1) by eliminating
the vent holes which are required when using a dry reagent
that requires diluent and (2) by adding an additional
sealing panel 144 which will be described in greater detail

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2 below. The cuvette 110 is comprised of a hollow body 112
3 (providing a rectangular parallelepiped interior chamber)
4 that includes side walls 118 uniformly molded as a single
5 unit with end walls 114 and a bottom wall 116 (Figure 9)
6 forming a hollow receptacle for receiving a liquid reagent,
7 prepackaged at the factory. The side walls 118 include an
8 optically clear window 118a on opposite side walls 118 to
9 permit the optical or ocular testing for the liquid reagent
10 alone and when subsequently mixed with the specimen. One
11 locating key 156 is centered on each side to locate the
12 cuvette with respect to the automated testing machine.

13 The cover 120 is ultrasonically or otherwise
14 bonded to the body 112 and includes a circular aperture 126
15 and a flat top wall 122 that includes a specimen chamber
16 formed by side walls 122a and bottom trap doors 136 which
17 are sealed together initially but are frangibly openable by
18 action of post 134 when the cap 128 is closed in the
19 automated test machine. A purge reservoir 124 for purging
20 the diluent dispenser tip in the automated equipment is
21 provided to wash out residual reagent and prevent carry over
22 into the next or following cuvette which may contain a
23 different reagent. The cover 120 includes a pair of arched
24 side members 138 which function and interact with the
25 automated machine as a roll down surface.

26 A sealing panel 144 is employed solely with a
27 cuvette using a liquid reagent, eliminating the need for a
28 diluent. The upper surface of top wall 122 also includes
29 grooves 154 longitudinally disposed along each edge which
30 receive glands 152 in the gasket/plug sealing panel 144 to
31 firmly seal panel 144 over the cover surface 122. The panel
32 144 also includes an aperture 150 which permits access to
33 the specimen chamber defined by walls 122a so that the
34 specimen can be added to the specimen chamber.

35 The cap 128 includes a modified post 134 having
36 inclined surfaces 134b terminating in knife-like edge end
37 tip 134a which acts as a plunger for piercing the specimen
38 chamber trap doors 136 at the centerline of the doors to

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2 ensure proper separation to effect direct hinge action of
3 the doors 136 to release the specimen into the cuvette body
4 112 containing the liquid reagent.

5 Referring now to Figure 9 the cuvette 110 is shown
6 from a bottom perspective view disclosing the bottom wall
7 116 having an optically clear window 116a which permits
8 fluorescent polarization and nephelometric type tests.

9 The cap 128 is unitarily connected by a hinge 130
10 which also acts to seal the end of the cuvette when the cap
11 128 is closed over the cover 120.

12 The cap 128 includes a pair of parallel ridges 160
13 which interact with the automated machine and rollers for
14 closing the cap 128 and resulting in a secure latching of
15 the cap.

16 A pair of snaps 132 which are integrally formed
17 with the cap 128 are engaged into slot edges 140a at the
18 opposite end of cover 120 to firmly attach the cap to the
19 cover when the cap is closed resulting in a leak-proof seal.

20 Referring back to Figure 8, in the manufacture of
21 the liquid reagent model, the plug/gasket sealing panel 144
22 is firmly attached at the factory to the top of cuvette
23 cover 120 after the liquid reagent has been inserted into
24 the cuvette body 112 chamber. With the panel 144 attached,
25 the liquid reagent is sealed tightly within the cuvette by
26 action of the panel 144 which includes plug 148 received
27 into aperture 126 and the fact that the specimen trap doors
28 136 in the specimen chamber 122a are also sealed.

29 To perform a test using the liquid reagent model,
30 the specimen is added to the specimen chamber 122a at the
31 testing site, which may be a doctor's office, laboratory or
32 the like. The cuvette 110 then is inserted into the
33 automatic testing machine in the cuvette's cap open position
34 as shown in Figure 8. The automated machine can first test
35 the liquid reagent alone in its present state (before the
36 specimen is mixed) while the cuvette 110 is located at a
37 test station in the automatic testing machine. After a first
38 measurement of the liquid reagent is obtained, optically or

1
2 ocularly, then the testing machine closes the cap 128,
3 causing the post 134 to fracture and open the two inclined
4 trap doors 136 causing the specimen to be received into the
5 interior chamber of cuvette body 112 containing the liquid
6 reagent. The cuvette is then boosted for mixing, causing a
7 rocking action on its radius corners adjacent the bottom
8 floor 116, greatly increasing the mix action. A second
9 measurement is then obtained optically or ocularly of the
10 liquid reagent-specimen mixture and the results calculated
11 by the testing machine.

12 A raised sealing bead 142a (Figure 8) is disposed
13 about the surface of cap surface 142. The bead is a raised
14 portion such that when cap 128 is closed firmly against
15 panel 144 (made of a silicone rubber), the bead 142a will be
16 impressed into the silicone rubber of panel 144, forming a
17 tight liquid impervious and air impervious seal about the
18 cover top surface 122 including the purge chamber 124.

19 Referring now to Figure 10, a modified cuvette
20 110a for use with a prepackaged dry reagent (powdered or
21 capsule) is shown which is inserted in the cuvette body 112.
22 Note that in this embodiment the plug/gasket panel 144
23 (Figure 8) is not used. The aperture 126 acts to receive a
24 diluent dispensing probe in the automated machine (not
25 shown) for dispensing diluent into the cuvette body 112 that
26 houses the dry reagent. The diluent is added at the test
27 site while performing the test in the automated machine. The
28 purge reservoir 124 collects diluent purged from the diluent
29 dispensing tip (not shown) after the dispensing tip has been
30 withdrawn from aperture 126 during the testing process. In
31 this alternate embodiment three vent slots 162, 164 and 166
32 are provided which collectively allow for air to escape from
33 the cuvette body 112 while the diluent is being added.

34 A thin frangible film 168 of hydrophobic metrical
35 polypropylene is used to cover the vent holes 162, 164 and
36 166 and diluent aperture 126, specimen chamber 122a and the
37 purge reservoir 124. The film 168 is gas permeable which
38 allows air to vent through the vent slots during diluent

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2 dispensing while also allowing easy fracture for insertion
3 of the specimen into the specimen chamber 122a. The film 168
4 provides another benefit in that the sealing beads 142a in
5 cap 128 will also engage the thin film when the cap 128 is
6 closed and locked over cover 120 again ensuring that the
7 cuvette is sealed after the test is completed. The film is
8 attached to the cover surface by a suitable glue.

9 In this embodiment the cuvette 110a functions
10 similarly to the original cuvette 10 discussed in Figures 1
11 through 7 with regard to the powdered and capsule reagents
12 in term of operation and the automated machine.

13 Figures 9 and 11 show another improvement in the
14 alternate embodiment of the invention that functions with a
15 carousel in the automated test machine which carries each
16 cuvette through the automated test machine. As shown in
17 Figures 9 and 11, the side end wall 114 includes, on each
18 end, a lateral, concave channel 114a which acts to receive
19 snap legs, which are in the automated test machine (not
20 shown) to firmly hold the cuvette in position relative to
21 the carousel and the machine. Also the lower corners 114b
22 formed by the side walls 114 and the bottom wall 116 have
23 equal predetermined radii of curvatures. At a particular
24 stage in the automated test machine, the curved corners 114b
25 engage surfaces having comparable curvatures on the snap
26 legs to produce a rocking motion whenever the carousel is
27 moved backward and forward to induce vigorous mixing within
28 the cuvette after the specimen has been introduced to the
29 reaction chamber and the cap is closed. Thus the curved end
30 portions 114b while engaging a similar surface of curvature
31 on the snap legs will permit a rocking motion back and forth
32 for improved mixing action. The bottom wall surface 116 is
33 substantially flat to permit a roller or wheel in the
34 automated test machine to raise the cuvette, disengaging the
35 snap legs from the snap leg detent 114a at the mixing stage
36 in the machine. Thus the cuvette is configured on its
37 outside and bottom walls to allow the snap legs in the
38 automated testing machine to accomplish the dual purposes of

1
2 allowing for rocking the cuvette for mixing while in the
3 other stages to secure the cuvette in position in the
4 carousel.

5 Figures 11 and 12 show the specimen receiving
6 chamber defined by walls 122a and bottom trap doors 136
7 which are sealed together along a frangible center line 136a
8 forming the bottom of the specimen chamber 122a.

9 The bottom trap doors 136 are inclined to reduce
10 possible trapping of bubbles when diluent or liquid reagent
11 are dispensed through aperture 126. In their initial state
12 the hinged doors 136 are joined together and to the walls
13 122a of the specimen chamber so that the compartment is
14 sealed from the body 112 chamber. The post 134 size and
15 shape is configured to more

16
17 easily fracture and open the trap doors 136 in the specimen
18 chamber when the cap 128 is closed tightly over the cover of
19 the cuvette. In the cross section as shown, the linear
20 weakened portions 136a and 136b between the walls 122a and
21 the doors 136 form an "H" shape. The trap doors 136 remain
22 sealed until forced open by the post 134 which causes the
23 centerline 136a and the wall edges 136b to frangibly detach
24 forcing the doors to open much like a pair of trap doors.
25 Note that the bottom floor is inclined relative to the
26 horizontal which aids in dispensing the specimen into the
27 reagent containing chamber 112. Note also from Figures 8 and
28 10 that the post 134 end tip includes a pair of inclined
29 surfaces 134b terminating in the edge 134a which meets the
30 centerline 136a of the trap doors 136 when the cap is closed
31 aiding the doors 136 to spread open from the center
32 position. This allows the specimen to be completely received
33 into the mixing chamber 112a where it mixes with either the
34 dry reagent and diluent mixture or with a liquid reagent.

35 Figures 11 and 12 also show hinge 130 that
36 connects the cap 128 (partially shown) unitarily to the
37 cover 120.

38 The present invention provides a disposable, non-

1
2 reuseable cuvette that allows a technician in a doctor's
3 office or laboratory to perform a variety of medical
4 diagnostic tests quickly, safely, accurately and
5 inexpensively by significantly reducing the technician
6 loading activities required and the structural complexity of
7 the cuvette and its interaction with the testing machine.

8 The instant invention has been shown and described
9 herein in what it is considered to be the most practical and
10 preferred embodiment. It is recognized, however, that
11 departures may be made therefrom within the scope of the
12 invention and that obvious modifications will occur to a
13 person skilled in the art.
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What I claim is:

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1. A disposable cuvette for performing a diagnostic test on a sample in an automated machine capable of measuring the radiant energy absorbed by a sample under analysis comprising:

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a hollow body forming a container with an open top;

a cover sealably attached over the open top of said body, said cover including a sample receiving chamber having a bottom floor, and said cover having a first aperture for the introduction of a diluent into said container body; and

a cap sized to fit over said cover to seal said first aperture and said cap including means for piercing the bottom floor of said sample receiving chamber when said cap is attached to said cover, sealing the cuvette.

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2. The cuvette as in claim 1, wherein:

the body container volume is predetermined to contain when full the required amount of diluent for a predetermined test to be performed in said cuvette.

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3. The cuvette as in claim 1, including:

a reagent disposed in said container body of predetermined kind and quantity for performing a particular test on a sample.

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4. A cuvette as in claim 1, wherein:

said sample receiving chamber bottom floor is a thin film that can readily be pierced by said means to pierce said sample chamber.

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5. A cuvette as in claim 1, wherein:

said sample receiving chamber bottom floor is a moveable closure that is activated to open when said cap is placed in a closed position relative to said cuvette.

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6. A cuvette as in claim 1, wherein:

said sample receiving chamber in said cover includes a

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2 thin film cover having an aperture therein for admitting the
3 end of a pipette.
4

5 7. A cuvette as in claim 1, including:
6 moveable flaps mounted over the cover first aperture
7 for admission of a diluent dispenser.
8

9 8. A cuvette as in claim 1, wherein:
10 said cap is integrally formed with said cover and said
11 cap hinges.
12

13 9. A cuvette as in claim 1, wherein:
14 said cover includes a pair of parallel, upwardly
15 disposed ridges for mechanical interface of said cuvette
16 within an automated testing machine.
17

18 10. A cuvette as in claim 1, wherein:
19 said cover having vent holes to aid in full diluent
20 dispensing and sized in diameter to prevent diluent passage
21 thereinto because of the surface tension of the diluent.
22

23 11. A cuvette as in claim 1, wherein:
24 said sample receiving chamber bottom floor is a thin
25 membrane having a breakline of minimum thickness formed in
26 the injection molding process that is pierced when said cap
27 is placed in a closed position relative to said cuvette
28 cover.
29

30 12. The method of performing automated, medical diagnostic
31 testing on a sample in a cuvette comprising the steps of:

32 a) providing a cuvette with a reaction chamber;

33 b) pre-loading said cuvette reaction chamber with a
34 particular reagent for performing a particular diagnostic
35 test;

36 c) loading the sample at the test site into a frangible
37 receptacle in said cuvette above said reaction chamber;

38 d) loading said cuvette into an automated testing

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machine;

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e) filling the reaction chamber with diluent;

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f) mixing the reagent and diluent in the reaction chamber;

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g) measuring the emergent radiation of the reagent-diluent mixture to obtain a baseline absorptivity measurement;

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h) sealing said reaction chamber while simultaneously rupturing said sample container receptacle causing said sample to be admitted to said reaction chamber;

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i) mixing said reagent, diluent and said sample; and

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j) measuring the emergent radiation of the contents of the reaction chamber.

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13. A cuvette for performing an automatic diagnostic test on a specimen comprising:

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a hollow body having a bottom wall and a plurality of side walls unitarily formed defining a rectangular reagent receiving chamber, at least two opposing side walls having windows sufficiently transparent to permit radiant energy transmission for optical or ocular testing;

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a substantially rectangular cover having a flat central portion and raised side ridges, said cover sealed to the upper end of said hollow body side walls, said cover including a specimen chamber having side walls extending downwardly from a specimen chamber aperture in said cover, said specimen chamber including a pair of movable closures forming the bottom floor of said specimen chamber, said movable closures initially joined together along opposing edges and including a frangible joint therebetween, each removable closure hingedly connected to said opposite specimen chamber side walls, said cover including an aperture for receiving a liquid dispensing probe for dispensing a liquid into said reagent receiving chamber;

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a sealing cap having an upper surface and a lower surface relative to a closed position for sealing said cover openings, said sealing cap hingeably connected to one end of

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2 said cover, said cap including an elongated post extending
3 substantially perpendicularly from the lower surface of said
4 cap and positioned and sized to engage and open the movable
5 closures forming the bottom wall in said specimen chamber
6 whenever said cap is in a closed position over said cover.
7

8 14. A cuvette as in claim 13, wherein:

9 said specimen chamber floor including said movable
10 closures is inclined angularly relative to the bottom wall
11 of said hollow body to reduce the possible trapping of
12 bubbles when diluent or liquid reagent are being dispensed
13 into said reagent chamber.
14

15 15. A cuvette as in claim 13, including:

16 a raised bead disposed in predetermined areas around
17 the periphery of the cap lower surface to aid in sealing the
18 cap to said cover when said cover is in a closed position.
19

20 16. A cuvette as in claim 13, wherein:

21 said cap post is configured to include an elongated
22 shaft body and a shaft tip, said shaft body having a cross-
23 shaped cross section and said shaft tip having a pair of
24 flat surfaces angularly inclined relative to each other to
25 form a knife-like edge substantially aligned with the
26 centerline between said movable closures of said specimen
27 chamber when said cap is closed.
28

29 17. A cuvette as in claim 13, wherein:

30 a purge reservoir formed in said cover near one end of
31 said cover, said purge reservoir including a plurality of
32 walls and a floor formed within the cover structure.
33

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35 18. A cuvette for use with a liquid reagent as in claim 17,
36 including:

37 a sealing gasket sized to fit over a predetermined
38 portion of said upper cover surface, said sealing gasket

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including a plug that is received into said diluent receiving aperture, and a specimen chamber aperture to permit access to said specimen chamber.

19. A cuvette for use with a dry reagent as in claim 17, including:

air venting means provided through said cover surface layer to vent said reagent chamber; and

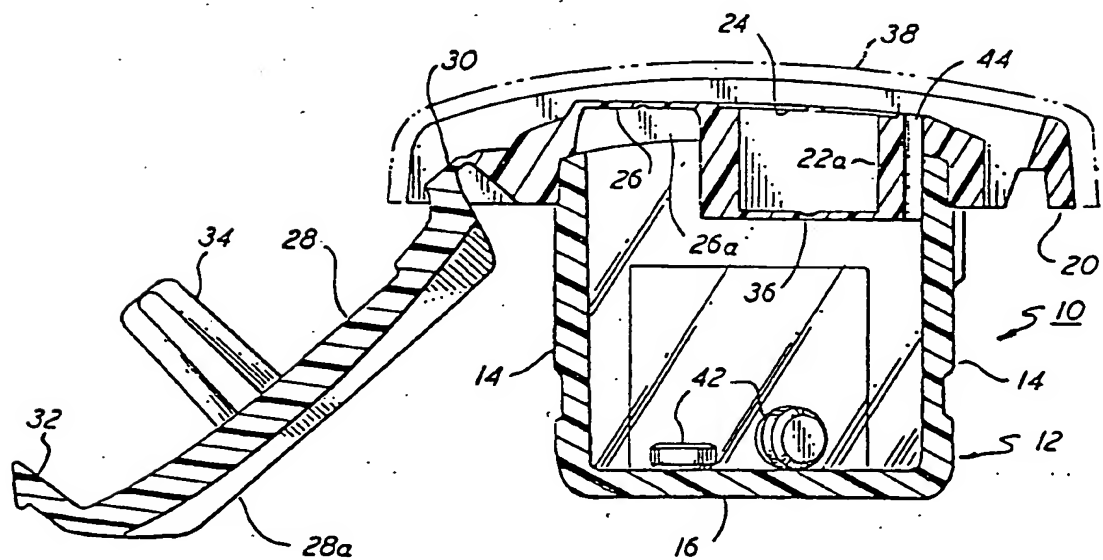
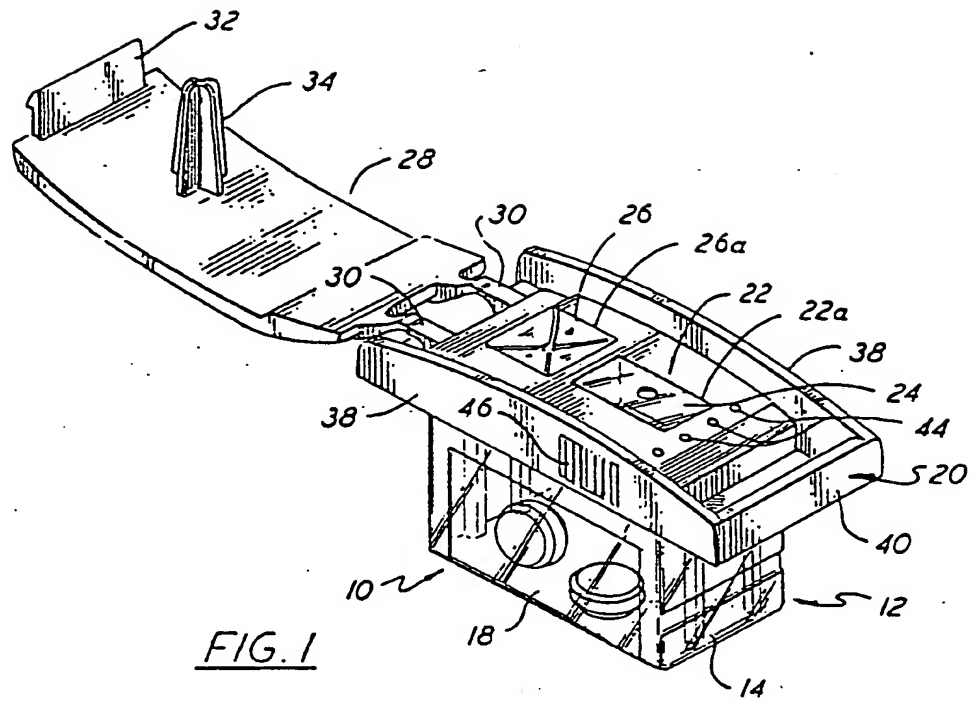
thin gas permeable film affixed to said flat central cover portion, overlaying said dispensing probe aperture, said specimen chamber aperture and said venting means.

20. A cuvette as in claim 17, wherein:

said hollow body bottom wall includes a window sufficiently transparent to permit transmission of radiant energy for fluorescent polarization and nephelometric testing.

21. A cuvette as in claim 17, wherein:

said hollow body side walls and bottom wall having exterior surfaces including detents strategically located for engagement with snap legs in said automated test machine and forming edges of a predetermined radius of curvature to permit vigorous rocking motion of said cuvette in said automated test machine.



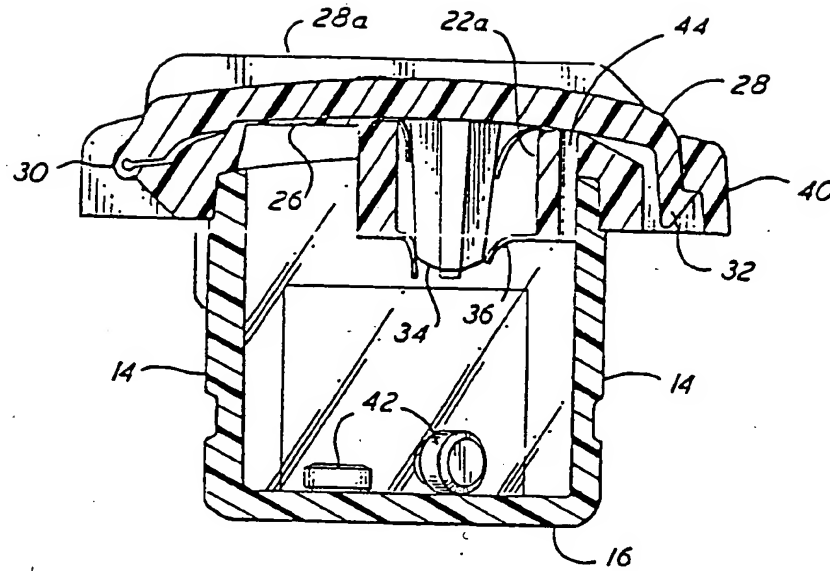


FIG. 3

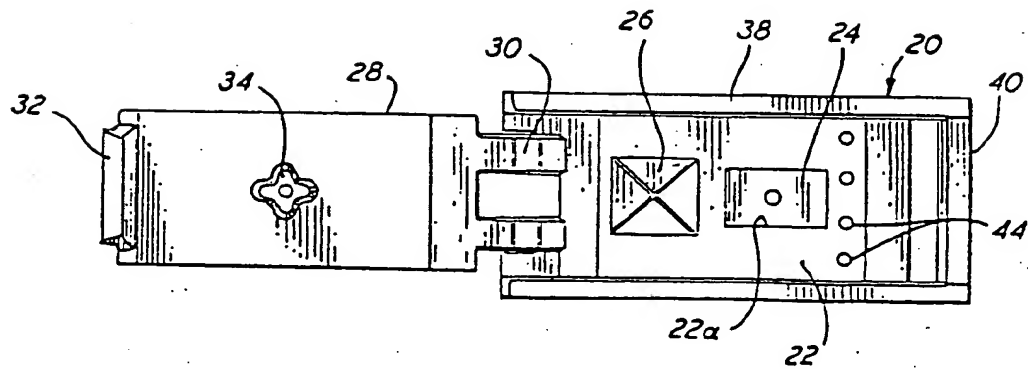
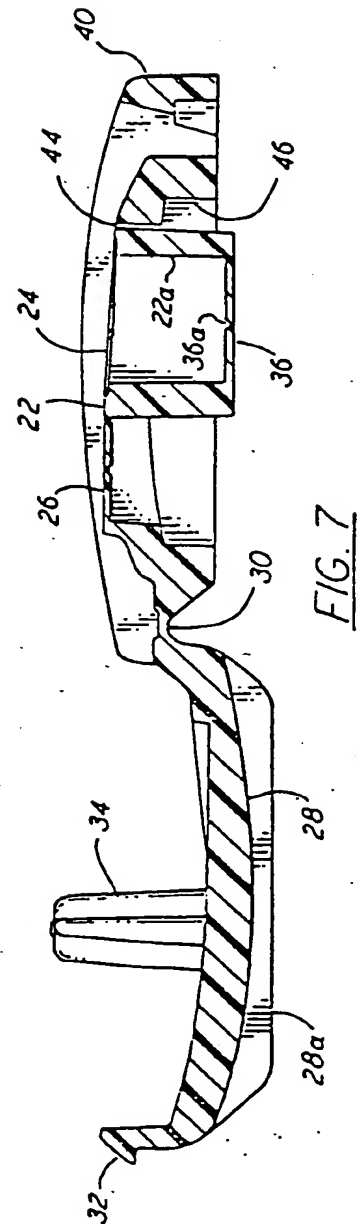
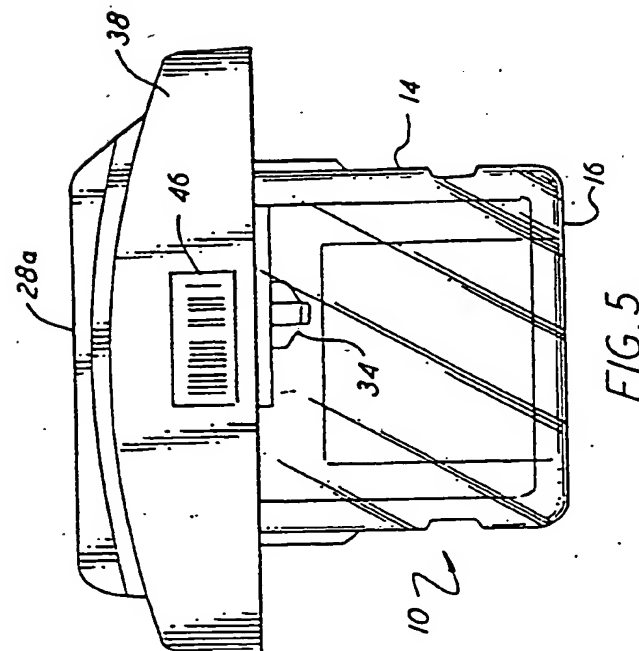
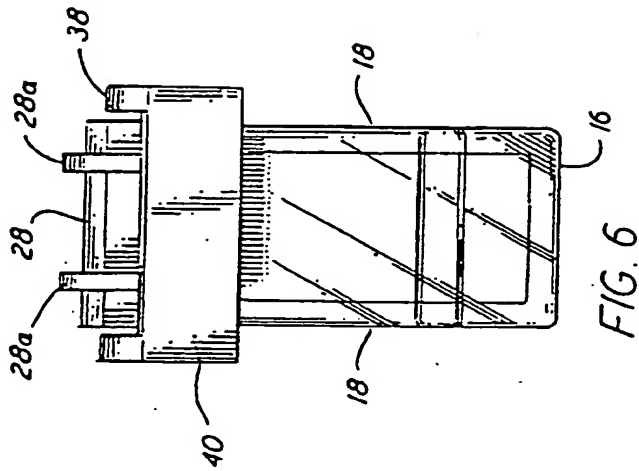


FIG. 4



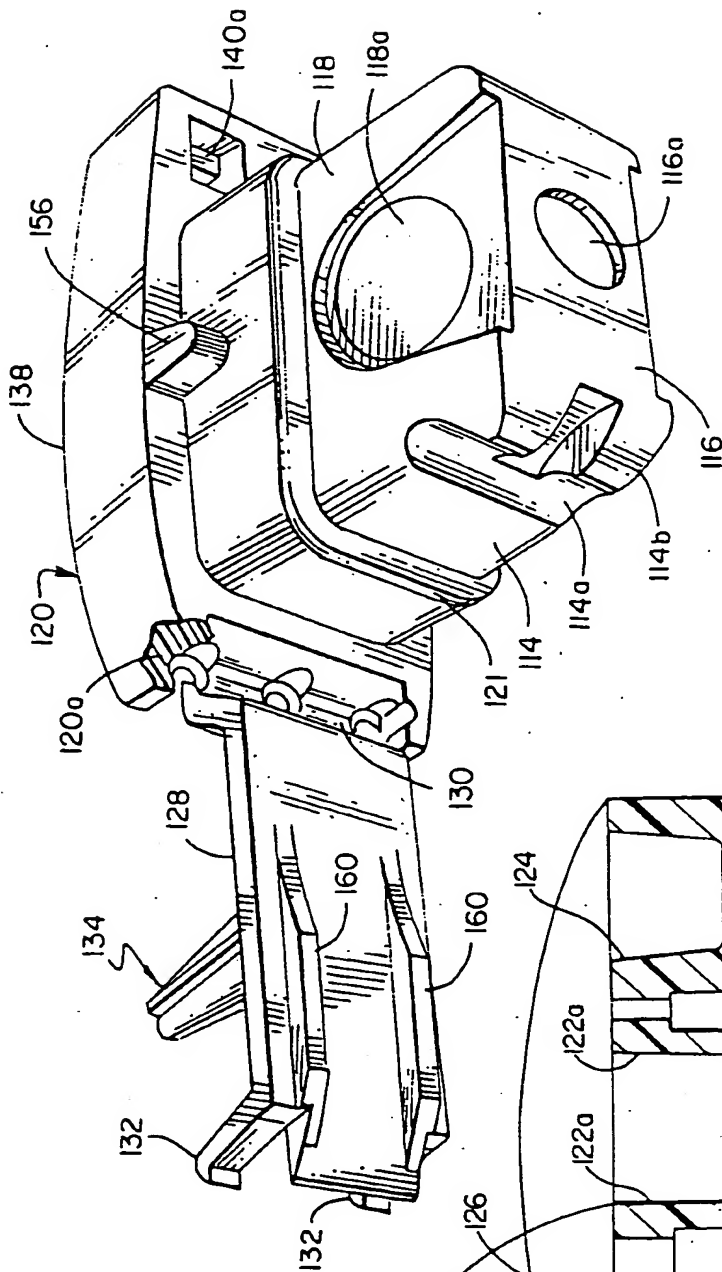


FIG. 9

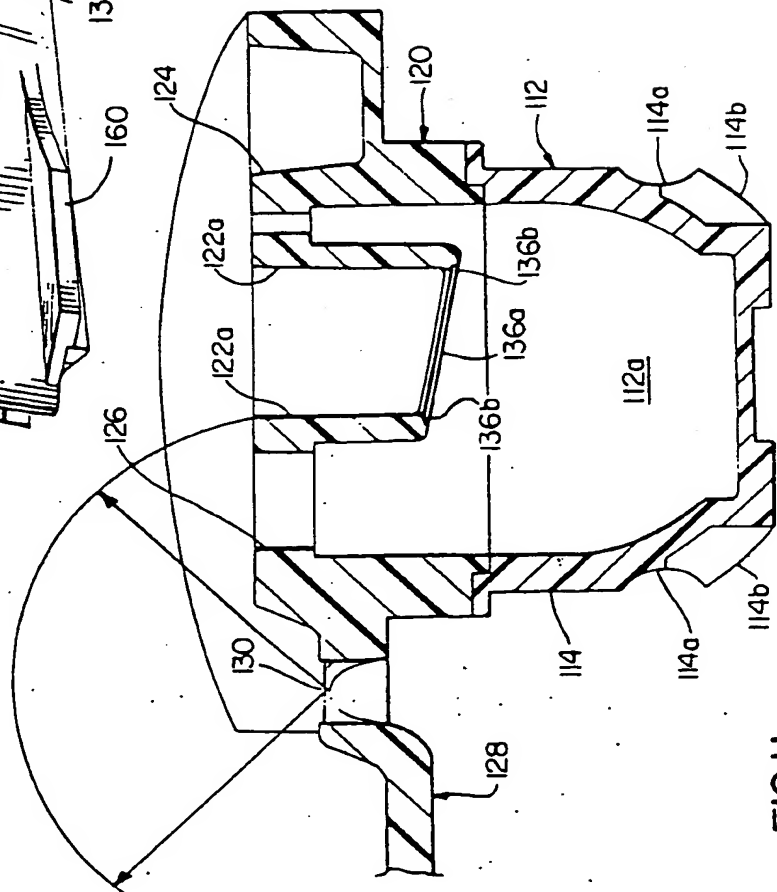


FIG. 11

